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DISTRICT COURT
    STATE OF MINNESOTA
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                                   NINTH JUDICIAL DISTRICT
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    COUNTY OF POLK
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                                    ) File No. 60-CR-06-8233
    State of Minnesota,
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                         Plaintiff, )
                                           TESTIMONY OF
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               VS.
                                           JOHN MARTIN
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    John Gerard Miller,
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                         Defendant.
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             The above-entitled matter came on for hearing
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    before the Honorable Tamara L. Yon, one of the judges of
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    the within court, in the courtroom in the courthouse at
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    Crookston, Minnesota, on September 29, 2010.
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                Mr. Greg Widseth, County Attorney in and for
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    said County of Polk, appeared on behalf of the State of
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    Minnesota.
                Mr. Eric Gudmundson, Assistant Public
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    Defender for the Ninth Judicial District, appeared on
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    behalf of the Defendant, who was also present in court.
                THEREUPON, the following proceedings were
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    had:
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JOHN MARTIN, 1 being first duly sworn, was examined and testified by 2 3 ITV as follows: THE COURT: Mr. Widseth. 4 DIRECT EXAMINATION 5 BY MR. WIDSETH: 6 Could you please state your full name for the 7 Court? 8 John Martin. J-o-h-n M-a-r-t-i-n. A. 9 Q. And what do you do for a living, Mr. Martin? 10 I'm a toxicologist. Specifically, my job duties 11 are technical consultant and certifying scientist for 12 13 Redwood Toxicology Laboratory. Q. And where is Redwood Toxicology Laboratory 14 15 located? A. In Santa Rosa, California. 16 Q. And as the toxicologist in certifying -- first of 17 all, how long have you been employed with Redwood 18 19 Toxicology? I've been with Redwood Toxicology for over 20 21 11 years. Q. And how long have you been a toxicologist? 22 I've been in the field of clinical laboratory 23 science specifically for drugs of abuse testing for over 24 30 years. 25

- Q. And do you have any professional degrees?
- A. Yes. I have a bachelor of arts in biology, a
 master of arts in biology, a master's of science in
 clinical laboratory science, and I'm also licensed under
 the Federal guidelines which are the Clinical Laboratory
 Improvement Act of 1988 as administered by the State of
 California and I have been so licensed for over
 - Q. And do you have any other licensures other than that one?
 - A. No. I'm also a member of the California
 Association of toxicologists.
 - Q. And in your employment with Redwood Toxicology have you been involved in the laboratory's use of EtG or EtS testing?
 - A. Yes.

30 years.

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- 17 | Q. How long have you been involved with that?
- A. Since the very beginning. We started research on testing back in March of 2006.
 - Q. And how long has Redwood Toxicology then used -- and could you tell the judge kind of what the EtG and EtS testing is generally?
- A. Certainly. There had been found and have been considered a time prior to that, I think it's like 1953, that when there was alcohol present in the liver, that

there would be -- it would be metabolized to certain products. Two of these products were ethyl glucuronide and ethyl sulfate. They are formed by different types of metabolism, so they are truly different metabolites of ethanol as detected in urine.

The analysis that's used -- there were a number of different analyses tried in the beginning, but today the highest level of testing is that by Liquid Chromatography/mass spectrometry/mass spectrometry or LC/MS/MS and it's that methodology that allows for quantitation at low levels of drugs. And the procedure that we developed is one that's published in the scientific literature and accepted there as a method for detecting EtG and EtS.

Basically what the protocol or standard that's used for testing is that a curve is run. In other words, a number of different samples of standards that are known and then after that is performed, controls are run. Those controls are at three different levels and also include a negative. And if those are acceptable, then the run can be processed at that time. And what it looks at is a number of different variables. It's often referred to when GCNS testing is used as finding a fingerprint in that each analyte has different parts of it that can be measured and those parts, as

they appear, have to match those of the standards so that the percentages of each one of the different types has to match as well as a specific time that it occurs, from the time the sample is interjected into the analyzer until the time it is at state. And so all of those things have to match that of the known standards before and have to match those in order to qualify to be called positive.

In our laboratory there have been recently newer methodologies that are starting to be used that are similar to the types of screens that are done routinely on urine, such that there may be sometime in the future screening processes that do not require the LC/MS/MS as the primary initial testing. In our laboratory we still use LC/MS/MS as the primary initial screening test. Once the sample goes through the initial screening process and it is determined to be positive, it is then taken and run a second time. That second time then allows for the quantitation of the specific analytes EtG and EtS. Once those are identified and those are verified by an analyst, then that run is reviewed by a certified scientist and then likewise, I have reviewed those results as well.

Q. So, just so the judge -- so we got the judge on the same page as we're on. The ethyl glucuronide -- if

you could pronounce that for me.

- A. Ethyl glucuronide, EtG.
- Q. EtG. And that's what I was going to get at.
- 4 That's EtG, right? And ethyl sulfate is the EtS?
 - A. Yes, sir.
 - Q. And those come as a metabolite I think you said of ingested ethanol then?
 - A. Yes.

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- Q. And how are they then detected using a urine sample? I mean, and I guess what I'm trying to get at is how do those metabolites get into the urine?
- A. Oh, the metabolism takes place in the liver and so if there is ethanol present and there are the specific metabolic pathways that then cause -- one is called glucuronidation and the other one is called sulfation. So one of the pathways then recreates the product EtG, whereas the second metabolic pathway creates the EtS or ethyl sulfate.
- Q. And that comes from ingested alcohol or ethanol you said then?
 - A. From ethanol, yes.
- Q. Is it possible, at least in your experience and your understanding, to have EtG or EtS metabolites appear in the urine through transdermal or touching of the skin with ethanol?

- A. Yes. There are a number of articles that are both published and unpublished that indicate that when there is the use of, as a class, hand sanitizers, some of those actually have as much as 62 percent ethanol. When there's the use of that product on the hands that there can be two different things that occur. There can be dermal and also there can be vapor. Most of the studies require the positive results obtained from the use of hand sanitizers to that of the ingestion of the vapors because it's so close.
- Q. And I might be getting ahead of myself here.

 Let's go back to one thing. If you take a urine sample from someone, how long can you determine the presence of alcohol within that urine sample?
- A. Okay, if we're talking about alcohol, alcohol itself is well-studied as far as under the influence and those vary depending on what type of ingestion there has been, but generally by the time ten or twelve hours have passed, the ethanol would be negative in the urine sample.
- Q. And when you use EtG or EtS testing or the LC/MS/MS to test for EtG and EtS, how long is the detection period then for ethanol?
- A. There is a longer window and that's one of the reasons for the advent of testing in alcohol programs in

that it also varies on the type of exposure that there may be and whether it's a single exposure or a multiple exposure and in cases where there truly was an abuse, those values can remain positive all the way up to 80 hours.

- Q. Okay. And is the use, I mean, you've got two different metabolites here. Does the presence of both of those have any greater significance as far as sensitivity or accuracy?
- A. Only in -- in all the initial studies EtG was referred to. And then only secondarily the testing became available and EtS then was tested. As I said, there are two different types of metabolism. There is a single literature citing that shows that if there are a number of criteria met that a person is diabetic, that they have used, that they have glucose and they form ethanol in their urine, that and the suppressants of specific micro-organisms, that there could be an increase in EtG due to all of those things being present. The same article also shows that the EtS, because it is a different metabolic pathway, is not affected the same. So in our laboratory we err on the side of negative in that we only report a positive test if both ethyl glucuronide and ethyl sulfate are used.
 - Q. And so I can get the judge at least on the terms

then, what I think she might be hearing is when I look back at the articles here, there seems to be two issues that come up with the testing for EtS and EtG and that's sensitivity and accuracy. Can you explain each of those to the judge?

- A. Well, certainly. The values have been found to be accurate, but the sensitivity is how low in a testing sequence you could go. And the original, all the original research tried to go to very low levels in that the test was used in order to monitor people in specific rehabilitation programs. And it was used as a counseling tool, so they wanted to be able to pick up essentially any, which was not possible, but extremely low levels and so that's one of the reasons that the cutoff levels were determined down to -- EtG down to 100 nanograms per mL and EtS down to 25 nanograms per mL.
- Q. Okay. So I at least get what I'm trying to get across to the judge here, these tests are very sensitive as far as it goes in determining the ingestion of alcohol, either through the skin or internally, is that correct?
- A. Yes.

Q. And then the issue of accuracy, at least as I've seen it, deals more with determining where that alcohol came from, would that be right?

- A. It can be that determination, yes.
- Q. Or the method of ingestion, I should say. So in this case you've seen the results or you were the certifying analyst on Mr. Miller's results, is that correct?
- 6 A. Yes.

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- Q. And do you have those results in front of you?
- 8 A. Yes, I do.
- 9 Q. And the test results were positive for both of these metabolites, is that correct?
 - A. Yes, they were.
 - Q. And the levels on the test result, which is Exhibit 1, under the EtG, there's a cutoff of 100 nanograms per milliliter, is that correct?
 - A. That's correct.
 - Q. And what does that represent to the Court, that cutoff level?
 - A. Essentially that's what I just mentioned. That cutoff level is the lowest level that the laboratory reports and it's the level at which values above that level then are often used in counseling situations.
- Q. And is that a common cutoff level within the drug testing industry?
- A. It is as far as the lowest level that's reported.

 The actual determination as far as what type of cutoff

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A. Yes.

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level to use for only the ingestion of ethanol and/or
secondary ingestion of ethanol has been -- a number of
different cutoff levels have been used since the time
the tests first started being performed.
  Q. And how long have these tests -- I mean how long
have we been testing for EtG?
  A. Our laboratory has been testing since March of
2006.
   Q. And when did it come to the United States then,
EtG testing?
  A. The first, in a research only, it was back in the
1950's.
   Q. Okay. So the cutoff level, as I understand it,
is kind of that minimal level where at least your lab
determines that further testing of that sample to
quantify the amount of EtS or EtG is done, is that
right?
   A. Yes.
   O. And then you do, as I think you indicated, the
second test and that's where you come out with the
actual numerical results for that urine sample?
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And you said there is a number of cutoff levels

that can be used at that point to determine

incidental -- I think the terms that I see are

incidental ingestion from intentional ingestion, is that correct?

- A. I said can be called incidental or secondary exposure.
- Q. And where does that level get set then in any particular testing regiment? Who sets that?
- A. There isn't anyone who specifically sets that.

 There have been recommendations made by the Substance

 Abuse and Mental Health Services Administration. There

 have been recommendations made by a number of the top

 researchers and there have been recommendations made by

 the U.S. Drug Court Professionals.
- Q. And can you give the judge kind of the gist of where all of those recommendations fall as far as where that cutoff level should be?
- A. Certainly. When the testing first began, it was thought that a cutoff level of 250 could be a reasonable cutoff level and these things changed over time and the studies started to begin to be available that showed that the use of some common products, cough medication, mouthwash, hand sanitizers, and even in some cases food products all contained ethanol, and when those were ingested, that they came up with levels of EtG.

And so then in reference then the cutoff levels were starting to be raised because in some of

these they showed that there could be, and most of the studies showed, not the hand sanitizers, but with the mouthwash and with the cough medicine that there could be levels as high as 250 with the ingestion of those products. And some of those products then were in excess, but not in excess to the amount that it would be equivalent to one or two drinks.

And so then there's also been studies on -in hand sanitizers as well and most of those studies
indicate that the levels are less than 100. Although,
there are a couple of studies where excessive use of the
product which was -- I think I gave you a copy of that,
but the excessive use they used a large quantity and it
was so much that it couldn't be just rubbed on the
hands. They had to run it all the way up and down their
arms and it actually hurt their eyes after time, but in
that case there was one sample that was as high as 700.
So even if we were using a 500 cutoff, then that would
indicate that there could have been exposure only due to
secondary.

However, in this case when we're dealing with this particular sample, there's a value that's considerably higher than 700. Some of the principal researchers have said that they may wish to use a cutoff level of a thousand or of 1,500, but those levels are

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hard to substantiate that there hadn't truly been the use of a secondary product in addition to the use of ingested ethanol, so it becomes difficult as far as interpreting.
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- Q. Are you aware of any scientific literature or papers regarding EtG or EtS testing which indicate that the incidental use in and of itself can result in EtG levels in excess of a thousand nanograms per milliliter? Are you aware of any generally accepted articles out there or peer review articles that would indicate that?
- A. There may be as far as specific researcher's findings with a single individual or something like that, but as far as those articles that have been reviewed, the highest levels that I've seen indicated are one was 713. There was another one that was in the 700s as well.
- Q. And for the Court's benefit, you had kind of put together a letter kind of summarizing your conclusions in these report results, is that correct?
 - A. Yes.
 - Q. And you sent a copy of that to me today?
- A. Yes.

- MR. WIDSETH: And, Your Honor, I would like to mark Exhibit 2, if I could, please.
 - THE COURT: You may.

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(Exhibit 2 was marked for identification.)
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                MR. WIDSETH: And I would offer Exhibit 2,
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    Your Honor.
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                MR. GUDMUNDSON: No objection.
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                THE COURT: Court will receive Exhibit 2.
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           (MR. WIDSETH CONTINUING) Now, Dr. Martin --
    Mr. Martin, with respect to the results that we have
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    with respect to Mr. Miller, is there anything further
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    that you can glean from the EtS result of 603?
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       A. Only in that similar to EtG there would be at
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    least a minimal cutoff of five times the lowest level
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    which would be 125, so it as well as the EtG are both
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    well above those secondary type cutoffs.
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       Q. And do these -- does EtG, does it peak at any
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    point? I assume like alcohol in the system it would go
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    on some sort of curve, is that right or is that
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    incorrect?
           No, that's correct.
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       A.
           And does the literature indicate that the EtG
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    numbers peak at any point?
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       Α.
           Yes.
           And at what point would that be?
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       0.
           Again, it may vary depending on the dose, but
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    generally with a one ounce of pure ethanol which would
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    be equivalent to one and a half to two drinks, the peak
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levels at four to eight hours could range anywhere from 700 to 2,000.

- Q. And do those peaks, do you know, do they hold true with respect to not only the intentional ingestion of alcohol but also the incidental circumstances or incidental exposures or secondary exposures as you call them?
- A. Each of the scientific articles do have some listing as far as when the peak levels were attained. And an example is with a Purell hand sanitizer. The one study found that the 713 was the highest and that was at nine hours. The other studies indicated similar in that eight to ten hours as far as a positive or peak value.
- Q. And did you, at least in looking at that issue regarding specifically here hand cleaners, did you see any of the scientific literature that's been peer reviewed or generally accepted out there which would indicate that you could have a result as high as Mr. Miller's here solely by the use of hand sanitizers alone?
- A. I haven't seen that article, but it doesn't mean that it would not be possible.
- Q. And at least in your professional opinion would this EtG level of 1130 and the EtS level of 603 on Mr. Miller's test, would that reflect incidental

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exposure to alcohol, in your opinion?

A. No. It could represent incidental alcohol in addition to ingestion of ethanol, but I don't believe it
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would be only secondary exposure.

- Q. So you don't believe that those results could come about only from the incidental or secondary type of exposure that we've been discussing?
 - A. Correct.

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- Q. It would also have to involve some form of at least intentional or ingestion of alcohol?
- 11 A. Additional exposure to alcohol, yes, ethanol.

MR. WIDSETH: I don't have any further questions at this time, Your Honor.

THE COURT: Mr. Gudmundson.

15 CROSS-EXAMINATION

16 BY MR. GUDMUNDSON:

- O. Good afternoon, Mr. Martin.
- 18 A. Good afternoon.
- Q. My name is Eric Gudmundson. I don't know if you recall talking to me on the phone sometime ago.
 - A. Oh, yes. Probably a month ago.
 - Q. Correct. I thank you very much for you being gracious with your time and you were very helpful to me. At that time one of the things I asked you was where I might go to find additional information on this subject,

do you recall that? 1 2 Α. Yes, sir. And I believe you directed me to possibly look up 3 Q. on the Internet a Dr. Gregory Skipper? A. Yes. 5 And you're familiar with him, is that right? 6 Yes. He's one of the first people that 7 investigated EtG in the United States. And you were aware that he maintained a website Q. 9 on the subject and so you directed me there, is that 10 11 correct? A. Yes. It's changed over time, but he does still 12 maintain a website, correct. 13 And you would, I assume, consider him an expert 14 0. 15 in this field? 16 A. Yes. Q. Now, one of the things that you testified about 17 was kind of a, at least at some point a disagreement 18 about an appropriate cutoff level, correct? 19 A. Yes. 20 Q. And you mentioned a couple of different 21 organizations or I guess I call them organizations. 22 I

don't know what they were, but can you list those again?

A. Well, certainly. There was an advisory by the

Substance Abuse and Mental Health Services

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Administration and it was, I think Dr. Skipper was one of the people on the panel as well as Paul Cary, but that, that advisory suggested against the use of EtG for punitive measures at low levels. There's a specific area in there, but it didn't go any further as far as delineating what would be a reasonable cutoff for secondary type exposures.

- Q. So you're familiar with that advisory?
- 9 A. Yes.

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- Q. And it hasn't been withdrawn, isn't that correct?
- 11 A. No.
- 12 Q. It has not been withdrawn, correct?
- 13 A. Correct.
 - Q. And you're aware that that advisory specifically indicated that the use of the EtG test in determining abstinence lacks sufficient proven specificity for use as primary or sole evidence that an individual prohibited from drinking, in a criminal justice or regulatory compliance context, has truly been drinking. You're aware of that, correct?
 - A. Yes.
 - Q. And that legal or disciplinary action based solely on a positive EtG test or other test discussed in this advisory is inappropriate and scientifically unsupportable, correct?